

14. A method according to claim 16, which comprises recovering the heptahelix receptor.

15. A method according to claim 16, wherein the host cell is a bacterium or yeast.

16. A heptahelix receptor having the sequence:

1 MNTTSSAAPPSSLGVEFISLLAIILLSVALAVGLPGNSFVV 40
41 WSILKRMQKRSVTALHVLNLALADLAVLLTAPFFLHFLAQ 80
81 GTWSFGLAGCRLCHYVCGVSMYASVLLITAMSLDRSLAVA 120
121 RPFVSQKLRTKAMARRVLAGIWLVSFLLATPVLAYRTVVP 160
161 WKTNMSLCFPRYPSEGHRAFLIFEAVTGFLLPFLAVVAS 200
201 YSDIGRRLQARRFRRSRRTGRLVVLILTFAAFWLPYHVV 240
241 NLAEARRALAGQAAGLGLVGKRLSLARNVLIALAFLSSSV 280
281 NPVLYACAGGGLLRSAGVGVFVAKLLEGTGSEASSTRRGGS 320
321 LGQTARSGPAALEPSPSESLTASSPLKLNELN (SEQ ID NO:2) 352

17. A fragment of heptahelix receptor comprising up to about 100 consecutive amino acid residues in Fig. 1 and containing Asn-2.

18. A fragment of heptahelix receptor comprising up to about 100 consecutive amino acid residues in Fig. 1 and containing Asn-164.

19. A fragment of heptahelix receptor comprising up to about 200 consecutive amino acid residues in Fig. 1 and containing Cys-90 and Cys-168.

20. A fragment of heptahelix receptor selected from the group consisting of DNA molecules having the sequences (I), (II), (III), (IV), (V), (VI), and (VII) in Fig. 1.

21. A method of detecting Burkitt's lymphoma, wherein the method comprises providing disrupted human cells, contacting the

cells with DNA as claimed in claim 1, and detecting a hybrid containing said DNA.

22. An antibody that specifically recognizes the heptahelix receptor as claimed in claim 16.

23. The antibody as claimed in claim 22, which is a monoclonal antibody.

24. A method of detecting Burkitt's lymphoma, wherein the method comprises providing human cells, contacting the cells with the antibody as claimed in claim 22, and detecting immunological complex containing said antibody.

25. A method for lowering the level of active leukotriene B4 in a mammal in need thereof, which comprises administering to said mammal a leukotriene B4-lowering amount of a receptor comprising the sequence of amino acids of SEQ ID NO:2.

26. A method for lowering the level of active leukotriene B4 in a mammal having inflammation, which comprises administering to said mammal a leukotriene B4-lowering amount of a leukotriene B4 receptor comprising the sequence of amino acids of SEQ ID NO:2.

27. A method for lowering the level of active leukotriene B4 in a mammal having bronchoconstriction, which comprises administering to said mammal a leukotriene B4-lowering amount of a leukotriene B4 receptor comprising the sequence of amino acids of SEQ ID NO:2.

28. A method for lowering the level of active leukotriene B₄ in a mammal having arthritis, which comprises administering to said mammal a leukotriene B₄-lowering amount of a leukotriene B₄ receptor comprising the sequence of amino acids of SEQ ID NO:2.

29. A method of treating a human to inhibit tissue injury accompanying inflammation resulting from leukocyte activity induced by LTB₄ produced in response to an inflammatory stimulus in the human, wherein the method comprises administering to said human a leukotriene B₄ receptor comprising the sequence of amino acids of SEQ ID NO:2 in an amount sufficient to inhibit activity of human leukotriene B₄ on polymorphonuclear leukocytes or monocytes in said human to thereby inhibit said tissue injury.

30. A method of treating a human to inhibit tissue injury accompanying inflammation resulting from leukocyte activity induced by LTB₄ produced in response to an inflammatory stimulus in the human, wherein the method comprises administering to said human a leukotriene B₄ receptor comprising the sequence of amino acids of SEQ ID NO:2 in an amount sufficient to modulate the inflammatory effect of leukotriene B₄ on polymorphonuclear leukocytes or monocytes by counteracting cell movement induced by LTB₄ in said human.

31. A method of treating a human to inhibit tissue injury accompanying inflammation resulting from leukocyte activity induced by LTB₄ produced in response to an inflammatory stimulus

in the human, wherein the method comprises administering to said human a leukotriene B₄ receptor comprising the sequence of amino acids of SEQ ID NO:2 in an amount sufficient to inhibit the stimulatory effect of leukotriene B₄ on adherence of polymorphonuclear leukocytes or monocytes in said human to thereby inhibit said tissue injury.

32. A method of treating a human to inhibit tissue injury accompanying inflammation resulting from leukocyte activity induced by LTB₄ produced in response to an inflammatory stimulus in the human, wherein the method comprises administering to said human a leukotriene B₄ receptor comprising the sequence of amino acids of SEQ ID NO:2 in an amount sufficient to inhibit stimulatory effect of leukotriene B₄ on oxidative burst of stimulated polymorphonuclear leukocytes in said human to thereby inhibit said tissue injury.

33. A method of treating a human to inhibit tissue injury accompanying inflammation resulting from leukocyte activity induced by LTB₄ produced in response to an inflammatory stimulus in the human, wherein the method comprises administering to said human a leukotriene B₄ receptor comprising the sequence of amino acids of SEQ ID NO:2 in an amount sufficient to inhibit the stimulatory effect of leukotriene B₄ on degranulation of stimulated polymorphonuclear leukocytes in said human to thereby inhibit said tissue injury.

34. A method of treating a human to inhibit tissue injury accompanying inflammation resulting from leukocyte activity induced by LTB₄ produced in response to an inflammatory stimulus in the human, wherein the method comprises administering to said human a leukotriene B₄ receptor comprising the sequence of amino acids of SEQ ID NO:2 in an amount sufficient to inhibit the effect of leukotriene B₄ on oxidative burst or degranulation of stimulated neutrophils in said human to thereby inhibit said tissue injury.

35. A method for assaying a ligand or an antagonist or agonist for said ligand, wherein the method comprises:

(A) providing a heptahelix receptor as claimed in claim 16 or a fragment thereof comprising a binding domain for the ligand, antagonist, or agonist;

(B) incubating the receptor with a test sample suspected to contain the ligand, antagonist, or agonist; and

(C) detecting binding between the receptor and the ligand, antagonist, or agonist.

36. A method according to claim 35, wherein the receptor is in an external cell membrane of a host cell transfected or transduced with DNA encoding the receptor.

37. A method according to claim 35, wherein binding is detected by intracellular calcium level in the host cell.

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